



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C12Q 1/28, G01N 33/72, 33/52	A2	(11) International Publication Number: WO 00/65083 (43) International Publication Date: 2 November 2000 (02.11.00)
(21) International Application Number: PCT/US00/11204 (22) International Filing Date: 26 April 2000 (26.04.00) (30) Priority Data: 09/299,399 26 April 1999 (26.04.99) US 09/517,481 2 March 2000 (02.03.00) US 09/517,441 2 March 2000 (02.03.00) US (71) Applicant (for all designated States except US): THE PROCTER & GAMBLE COMPANY [US/US]; One Procter & Gamble Plaza, Cincinnati, OH 45202 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): CAPRI, Maria, Grazia [IT/IT]; Via P. Nenni, 150, I-66020 S. Giovanni Teatino (IT). CARLUCCI, Giovanni [IT/IT]; Via A. Fieramosca, 118, I-66100 Chieti (IT). GUERRESCHI, Lisa [IT/IT]; Via G. Garibaldi, 25, I-26048 Sospiro (IT). HAMMONS, John, L. [US/US]; 7379 Dust Commander Court, Hamilton, OH 45011 (US). (74) Agents: REED, T. David et al.; The Procter & Gamble Company, 5299 Spring Grove Avenue, Cincinnati, OH 45217-1087 (US).		(81) Designated States: AE, AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), DM, EE, EE (Utility model), ES, FI, FI (Utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: FEMININE SANITARY DISPOSABLE ARTICLE HAVING A BLOOD DETECTION MEANS AS SENSOR		
(57) Abstract The present invention relates to feminine sanitary disposable absorbent articles and, more particularly, to sanitary napkins or panty liners comprising a blood detection means for menstruation prediction by having sensors which have a recognition element to detect the presence of blood indicative of the approach of menstruation and which provide a visual signal indicating the approach of menstruation.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

FEMININE SANITARY DISPOSABLE ARTICLE HAVING A BLOOD DETECTION MEANS AS SENSOR

FIELD OF THE INVENTION

The present invention relates to feminine sanitary disposable absorbent articles and, more particularly, to sanitary napkins or panty liners comprising a blood detection means for menstruation prediction by having sensors which have a recognition element to detect the presence of blood indicative of the approach of menstruation and which provide a visual signal indicating the approach of menstruation.

BACKGROUND OF THE INVENTION

Today, disposable articles, such as diapers, adult incontinence briefs, sanitary napkins, panty liners, interlabial devices and tampons, are widely used in feminine protection, infant and toddler care and in the care of incontinent adults as a means of containing, isolating and disposing of bodily wastes. These articles have generally replaced reusable, washable cloth garments as the preferred means for these applications because of their convenience and reliability. The disposable articles respond to a discharge event by absorbing or containing bodily wastes deposited on the article. Some disposable articles comprise a chemically reactive means to detect and signal various substances in the wearer's waste(s). However, none of these specifically detect or predict when a menstruation-related discharge event is about to occur and signal to the wearer or caregiver that occurrence.

The so far un-met consumer need of women which is targeted by the present invention is the desire to gain more information about the expected timing of their menstruation. More generally it is well established that women would like to know more about and monitor their bodily condition as relates to the status of their menstruation cycle as well as a pre-warning to any kind of related disorder advancing.

Therefore, the present invention contains two aspects which may have to be considered in a different claim set according to local patent law. One aspect is the diagnostic purpose in a medical sense of the development of the present invention while the other is the purely non-medical benefit of providing early information about the approach of menstruation in order to allow respective action by the consumer such as starting to use sanitary napkins or tampons.

A lot of disclosure exists on general diagnostic aspects relating to female menstruation cycle and genital/urinary tract related information. However, none of the references cited in the following does provide specific teaching to the present invention.

US 5,217,444 to Schönfeld et. al., published June 8, 1993 discloses absorbent pads such as tampons comprising a pH indicator material to indicate the acidity or alkalinity of a liquid by a color change. The pH indicator is intended to be wetted by vaginal secretions which are absorbed by the pad. Depending on the pH this will provide an indication of the health condition of the women's vaginal tract.

US 5,468,236 to Everhart et. al., published November 21, 1995 discloses a disposable absorbent product incorporating a chemically reactive substance which can provide visual indication of a chemical component in absorbed liquid such as vaginal discharges. The patent is un-specific as to its particular purpose

but exemplifies a disposable diaper in which a glucose indicating gel can be applied to the topsheet or the absorbent core. In a second example the chemical compound to be detected is a nitrate as indicator of urinary tract infections by GRAM negative bacteria. In particular Everhart provides for a reaction which is endpoint stable due to a catalytic sequence such that further liquid absorption will not alter the indication.

US 4,231,370 to Morz et. al., published November 4, 1980 discloses disposable diapers with a pH sensitive wetness indicator in a solid adhesive matrix.

US 5,823,953 to Richards et. al., published October 20, 1998 relates to a self diagnostic system for yeast or non-yeast related vaginal infections in the form of a catamenial pad or panty liner with detachable, color indicating pH strip mounted on the topsheet. The pH indicator identifies if the pH of the absorbed liquid is above or below a threshold value of 4.5 and is indented to allow health care decisions without a physician in respect to treating a yeast or a non-yeast vaginal infection.

WO 97/43955 to Buck et. al., published November 27, 1997 discloses a kit for home use for collecting vaginal fluid and exfoliated vaginal cells in menstrual fluid for diagnostic purposes. The fluid or cells are collected in an absorbent interlabial pad or in a pad which is placed inside an apertured housing which is placed inter vaginally for collecting fluid. The fluid can afterwards be used for diagnostic purposes as desirable.

EP 704 195, to Echeveria, published April 3, 1996 discloses a menstrual detector which comprises a sanitary napkin or panty liner including a chemical compound which provides a cooling reaction upon liquid absorption as indicator of the start of menstruation in order to allow the wearer sufficient time to replace

the indicating absorbent article by a full size and full capacity sanitary napkin or tampon.

Of course, many indicators as such are known and are usually independent of the source of the liquid of which analysis is desired. For example, EP 386 562 to Ismael et. al., published September 12, 1990 discloses a dry enzyme test composition with a color indicator result. Similarly, WO 90/06511 to Buck et. al., published June 14, 1990 discloses a stabilized indicator to determine the presence of an analyte in a liquid sample. Similarly, WO 89/11643 to Bouse et al., published November 30, 1989 discloses an indicator composition designed to increase the range of analyte concentration which can be analyzed.

EP 124 215 and EP 124 214 both to Oksman et al, published November 7, 1984 relate to wipe indicators for occult blood particular useful to analyze the presence of blood in feces. Similarly, EP 93595, to Wells, published November 9, 1983 relates to a dry diagnostic aid for use in an one-step determination of hemoglobin, especially in occult blood in feces.

EP 113 896 to Rothe, published July 25, 1984 relates to a chemical testing strip in which a test reagent is provided on a plastic film strip for quantitative evaluation. This patent exemplifies blood and glucose testing.

EP 101 980 and EP 101 979, both to Roy, published March 7, 1984 relate to the determination of dehydrogenase for the purpose of rapid clinical analysis and diagnosis. In the later case it is said that the carrier can be a tampon containing cellulose fibers.

In addition many publications in the medical field relating to pathological conditions of the female genital and urinary organs including methods of detecting and healing them have been published.

SUMMARY OF THE INVENTION

The present invention relates to absorbent sanitary articles for women which provide a visual blood detection of a quantity of blood in vaginal discharges deposited on the article which is not discernible by normal visual inspection by the human eye. The article is conventionally of a layered construction, each layer having a wearer and a garment facing surface. In general the article comprises a liquid permeable topsheet on the wearer facing surface, a liquid barrier backsheet on the garment facing surface, an absorbent core sandwiched between the topsheet and the backsheet. According to the present invention the article is preferably a sanitary napkin or panty liner, especially a thin panty liner with less than 3 mm thickness and most preferred a panty liner which is suitable to be worn in a string undergarment and has a generally triangular shape (also referred to as thong shape or thong liner respectively).

According to the present invention the article comprises a blood detection means which comprises a chemical composition of at least a first component selected from a peroxide or a per-acid or combinations thereof, and a second component selected from Gum Guaiac or tetra-methyl-benzidine or combinations thereof. The composition preferably further comprises a tensioactive material such as a surfactant or preferably a polyethylene-glycol.

In a preferred embodiment according to the present invention the first component is epta-phthalimido-peroxy-hexanoic acid and the second component is Gum Guaiac. The first and second components are preferably separated in the absorbent article, more preferably one component is provided on the garment facing surface of one layer and the other component is provided on the garment facing surface of a second layer or one component is provided on the wearer facing surface of one layer, preferably the core and the other component is provided on the garment facing surface of a the same layer.

When applying the composition to the article it is preferred to provide it in a liquid form, more preferably as a solution in an organic carrier which may evaporate after the composition has been applied. Application can be done by coating or preferably by spraying.

It is within the scope of the present invention to supplement the blood detector by additional menstruation related detectors such as additional biochemical or electro-chemical or chemical detection means suitable to detect physiological changes in the absorbed vaginal discharges which are indicative of the approach of menstruation and preferably create a separate, strengthened or modified visual signal to the user of such articles. Preferably such an additional detection means comprises a pH indicator or a progesterone hormone or estrogen hormone indicator or combinations thereof.

DETAILED DESCRIPTION OF THE INVENTION

DEFINITIONS

As used herein, the term "absorbent article" refers to devices which absorb and contain body exudates, and more specifically, refers to devices which are placed against or in proximity to the body of the wearer to absorb and contain the various exudates discharged from the body. The term "disposable" is used herein to describe absorbent articles which generally are not intended to be laundered or otherwise restored or reused as an absorbent article (i.e., they are intended to be discarded after a single use). As used herein, the term "joined" encompasses configurations whereby an element is directly secured to another element by affixing the element directly to the other element, and configurations whereby an element is indirectly secured to another element by affixing the element to intermediate member(s) which in turn are affixed to the other element. A "unitary" absorbent article refers to absorbent articles which are formed of separate parts united together to form a coordinated entity so that they do not require separate

manipulative parts like a separate holder and liner. A preferred embodiment of an absorbent article of the present invention is a unitary disposable sanitary napkin or panty liner or interlabial absorbent article, however also disposable incontinence briefs, incontinence undergarments, absorbent inserts, and tampons can benefit from the present invention if used during the period prior to menstruation.

The absorbent article, according to the present invention is conventionally constructed of three main elements: The topsheet, facing the user of the article during use and being liquid pervious in order to allow liquids to pass into the article; the backsheet, providing a liquid containment such that absorbed liquid does not leak through the article, this backsheet conventionally provides the garment facing surface of the article; and the absorbent core sandwiched between the topsheet and the backsheet and providing the absorbent capacity of the article to acquire and retain liquid which has entered the article through the topsheet.

Many absorbent articles and constructions are known in the art and have been described in ample detail over time. All of such materials are useful in the context of the present invention, provided they do not interfere with the selected menstruation prediction sensors. In the following only those examples, which are particularly beneficial for the use in preferred absorbent articles according to the present invention are mentioned. Those skilled in the art will readily be able to identify alternative materials which can also be used and which maybe particularly desirable in the context of menstruation predicting absorbent articles according to the present invention.

Topsheet

In general the topsheet is compliant, flexible, soft feeling and non-irritating to the wearer's skin. The topsheet preferably also can have elastic characteristics

allowing it to stretch in one or two directions. As used herein, the term "flexible" refers to materials that are compliant and will readily conform to the general shape and contours of the human body.

The topsheet can be made from a nonwoven or woven material or a film which has been rendered liquid pervious by aperturing. The topsheet can also be provided as a composite material or be comprised of more than a single layer, e.g. it can have a secondary topsheet or flow control layer. Such films and nonwovens or wovens can be made for example from polymers such as polyethylene or polypropylene compositions. Conventionally such polymers have been filled with coloring material such as titanium dioxide to provide a white opacity. Such a topsheet can also be provided without the coloring material to provide a transparent film or nonwoven which has a high degree of light transmission while the absence of a color filler, such as titanium dioxide, does not cause any substantial material changes. Such a topsheet would be particularly beneficial in the context of menstruation predictors which are inside the article to allow visual inspection whether a color change has happened. Hence the absence (or reduction below 15%, i.e. about half of the conventional quantity) of an opaque color filler in the polymeric material from which the topsheet can be made provides the additional benefit of better inspection characteristics (besides less costs and a reduced environment burden). It would also be possible to provide a non opaque, tinting color filler which allows transparency or partial transparency while providing a color. This tinting may beneficially be used to hide the initial or final color of the color indicators according to the present invention, and is preferably pale yellow or blue.

The topsheet can be transparent or can be only partially transparent. It is however sufficient for most embodiments according to the present invention if the topsheet is translucent to allow visual inspection of a color change.

Backsheet

In general the backsheet is compliant, flexible and soft feeling. The backsheet prevents the exudes absorbed and contained in the absorbent core from wetting clothes that contact the absorbent article such as undergarments. Preferably the backsheet is impervious to liquids (e.g., menses, sweat and/or urine). It can be manufactured from a thin plastic film, although other flexible liquid impervious materials can also be used. As used herein, the term "flexible" refers to materials that are compliant and will readily conform to the general shape and contours of the human body. The backsheet preferably also can have elastic characteristics allowing it to stretch in one or two directions.

The backsheet can comprise a woven or nonwoven material, polymeric films such as thermoplastic films of polyethylene or polypropylene, or composite materials such as a film-coated nonwoven material or fiber coated film. Conventionally absorbent articles comprise a backsheet of a polyethylene film having a thickness of from about 0.012 mm to about 0.051 mm, which can be impervious or rendered micro-porous by use of an inert filler and subsequent mechanical stretching.

The backsheet is preferably breathable, i.e. allows the transmission of water vapor, or even more preferable the transmission of air, however without sacrificing its main purpose to provide leakage protection for absorbed liquids. The backsheet can also comprise more than one breathable layer so as to replace a single breathable backsheet layer by at least 2 or 3 layers of a different or the same material. In particular two breathable layers, in which e.g. the one providing the wearer facing surface is a formed, apertured film with a three dimensional structure and e.g. the other, garment facing layer is a non-woven composite of melt-blown and spun-bonded fibers, are preferred breathable backsheet constructions. As with the topsheet the backsheet can be provided from a transparent or translucent material which would allow easier inspection of

detectors inside the article. Similarly it can also be provided with a tinted color filler, preferably pale yellow or blue.

Absorbent core

Conventionally the absorbent core can be a single entity or comprise several layers. It can include the following components: (a) optionally a primary fluid distribution layer; (b) optionally a secondary fluid distribution layer; (c) a fluid storage layer; (d) optionally a fibrous layer underlying the storage layer; and (e) other optional components.

a. Primary Fluid Distribution Layer

One optional component of the absorbent core according to the present invention is the primary fluid distribution layer. This primary distribution layer typically underlies the topsheet (if present) and is in fluid communication therewith. The primary distribution layer acquires body fluid for ultimate distribution to the storage layer. This transfer of fluid through the primary distribution layer occurs mainly in the thickness, but may also provide distribution along the longitudinal and transverse directions of the thong liner.

b. Optional Secondary Fluid Distribution Layer

Also optional according to the present invention is a secondary fluid distribution layer. This secondary distribution layer typically underlies the primary distribution layer and is in fluid communication therewith. The purpose of this secondary distribution layer is to readily acquire bodily fluid from the primary distribution layer and distribute it along the longitudinal and transverse directions of the thong liner before transfer to the underlying storage layer. This helps the fluid capacity of the underlying storage layer to be fully utilized.

c. Fluid Storage Layer

Positioned in fluid communication with, and typically underlying the primary or secondary distribution layers, is a fluid storage layer. It comprises preferably but not necessarily super-absorbent gelling materials usually referred to as "hydrogels," "superabsorbent" "hydrocolloid" materials. Absorbent gelling materials are those materials that, upon contact with aqueous fluids, especially body fluids, imbibes such fluids and thus form hydrogels. These absorbent gelling materials are typically capable of absorbing large quantities of aqueous body fluids, and are further capable of retaining such absorbed fluids under moderate pressures. In the prior art these absorbent gelling materials are typically in a granular form of discrete, non-fibrous particles. However, according to the present invention these super-absorbent gelling materials can also be provided in non-granular form, preferably in a fibrous form. If no absorbent gelling materials are provided then the storage layer can be provided by the material conventionally used as carrier material disclosed below.

In the fluid storage layer these absorbent gelling materials can be dispersed homogeneously or non-homogeneously in a suitable fibrous matrix also referred to as carrier. Suitable carriers include cellulose fibers, in the form of fluff or tissues, such as is conventionally utilized in absorbent cores. Modified cellulose fibers such as the stiffened cellulose fibers or viscose fibers can also be used. Synthetic fibers can also be used and include those made of cellulose acetate, polyvinyl fluoride, polyvinylidene chloride, acrylics (such as Orlon), polyvinyl acetate, non-soluble polyvinyl alcohol, polyethylene, polypropylene, polyamides (such as nylon), polyesters, bi-component fibers, tri-component fibers, mixtures thereof and the like. Preferred synthetic and man-made fibers have a denier of from about 3 denier per filament to about 25 denier per filament, more preferably from about 5 denier per filament to about 16 denier per filament. The carrier fibers can be provided as carded, spun-bonded, melt-blown, wet-laid, air-laid

substrates or combination of such lay down methods or combinations of such substrates.

Also preferably, the fiber surfaces are hydrophilic or are treated to be hydrophilic. A storage layer can also include filler materials, such as Perlite, diatomaceous earth, Vermiculite, etc., that lower rewet problems. Further the storage layer may comprise a binder including but not limited to Latex binders which can be sprayed as an aqueous solution onto the surface of the storage layer prior to curing.

If the absorbent gelling materials are dispersed non-homogeneously in a fibrous matrix, the storage layer can be locally homogeneous, i.e. have a distribution gradient in one or several directions within the dimensions of the storage layer. Non-homogeneous distribution thus includes e.g. laminates of the fibrous carriers enclosing the absorbent gelling materials or regions in which the absorbent gelling material has a different concentration relative to other regions.

If absorbent gelling material is present, the storage layer preferably comprises from 5% to 95% absorbent gelling materials, preferably from 5% to 50%, most preferably from 8% to 35%, absorbent gelling materials. Further the storage layer can comprise from 5% to 95% carrier fibers, preferably from 95% to 50%, most preferably from 92% to 65% carrier fibers.

Suitable absorbent gelling materials for use herein will most often comprise a substantially water-insoluble, slightly crosslinked, partially neutralized, polymeric gelling material. This material forms a hydrogel upon contact with water. Such polymer materials can be prepared from polymerizable, unsaturated, acid-containing monomers. Suitable unsaturated acidic monomers for use in preparing the polymeric absorbent gelling material used in this invention include those listed in U.S. Patent 4,654,039 (Brandt et al), issued March 31, 1987, and reissued as RE 32,649 on April 19, 1988. Preferred monomers include acrylic

acid, methacrylic acid, and 2-acrylamido-2-methyl propane sulfonic acid. Acrylic acid itself is especially preferred for preparation of the super-absorbent material, it also has a 'natural' transparency which is not optimal but acceptable if the desired transparency is not too high.

Whatever the nature of the basic polymer components of the hydrogel-forming polymeric absorbent gelling materials, such materials will in general be slightly crosslinked. Crosslinking serves to render the hydrogel-forming polymer gelling materials substantially water-insoluble, and cross-linking thus in part determines the gel volume and extractable polymer characteristics of the hydrogels formed from these polymeric gelling materials. Suitable crosslinking agents are well known in the art and include, for example, those described in greater detail in U.S. Patent 4,076,663 (Masuda et al), issued February 28, 1978. Preferred crosslinking agents are the di- or polyesters of unsaturated mono- or polycarboxylic acids with polyols, the bisacrylamides and the di- or triallyl amines. Other preferred crosslinking agents are N,N'-methylenebisacrylamide, trimethylol propane triacrylate and triallyl amine. The crosslinking agent can generally constitute from about 0.001 mole percent to 5 mole percent of the resulting hydrogel-forming polymer material. More preferably, the crosslinking agent will constitute from about 0.01 mole percent to 3 mole percent of the hydrogel-forming polymeric gelling material.

The slightly crosslinked, hydrogel-forming polymeric gelling materials are generally employed in their partially neutralized form. For purposes of the present invention, such materials are considered partially neutralized when at least 25 mole per-cent, and preferably at least 50 mole percent of monomers used to form the polymer are acid group-containing monomers that have been neutralized with a salt-forming cation. Suitable salt-forming cations include alkali metal, ammonium, substituted ammonium and amines. This percentage of the total monomers utilized which are neutralized acid group-containing monomers is referred to herein as the "degree of neutralization."

While these absorbent gelling materials have typically been disclosed in the prior art in granular form, it is possible in the context of the present invention that the absorbent gelling material is in a non-granular form for example as macrostructures such as fibers, sheets or strips or in a form suitable for coating, e.g. before curing or as part of an adhesive substance. A macrostructure can be prepared by forming the particulate absorbent gelling material into an aggregate, treating the aggregated material with a suitable crosslinking agent, compacting the treated aggregate to densify it and form a coherent mass, and then curing the compacted aggregate to cause the crosslinking agent to react with the particulate absorbent gelling material to form a composite, porous absorbent macrostructure. Such porous, absorbent macrostructures are disclosed, for example, in U.S. Patent 5,102,597 (Roe et al), issued April 7, 1992.

d. Optional Fibrous Layer

An optional component for inclusion in the absorbent cores according to the present invention is a fibrous layer adjacent to, and typically underlying the storage layer. This underlying fibrous layer would typically provide the same function as the secondary fluid distribution layer.

e. Other Optional Components

The absorbent cores according to the present invention can include other optional components normally present in absorbent webs. For example, a reinforcing scrim can be positioned within the respective layers, or between the respective layers, of the absorbent cores. Such reinforcing scrims should be of such configuration as to not form interfacial barriers to fluid transfer, especially if positioned between the respective layers of the absorbent core. Given the structural integrity that usually occurs as a result of thermal bonding, reinforcing

scrims are usually not required for the absorbent structures according to the present invention.

Another component which can be included in the absorbent core according to the invention and preferably is provided close to or as part of the primary or secondary fluid distribution layer are odor control agents. Typically active carbon coated with or in addition to other odor control agents, in particular suitable zeolite, silica or clay materials, are optionally incorporated in the absorbent core.

Physical characteristics of absorbent cores

Absorbent cores are usually non extensible and non-elastic, however, they can be rendered extensible and depending on the selected materials can also be made to have elastic characteristics. The term "extensible" as used hereinafter refers to a structure which under external forces such as those occurring during use extends in the direction of the forces or in the direction of a component of the forces in cases where only mono directional extensibility is provided.

The term "elastic" as used hereinafter refers to extensible structures which return at least partially to their initial state after the forces causing the extension cease to be exerted. Absorbent cores can be corrugated or pleated in one or several directions to provide a certain extensibility while selection of elastic fibers for the structure can provide elasticity.

The absorbent cores should preferably be thin. A thickness of less than 5 mm, preferably less than 3 mm and even more preferable between 0.8 and 1.8 mm is desirable such that the resulting articles can also have a low thickness. As with the topsheet or the backsheet transparency of or tinting of the materials forming the core can be particularly desirable to allow easy inspection of the detector status.

Non-limiting examples of panty liners and sanitary napkins which may be provided with a detection means include those manufactured by The Procter & Gamble Company of Cincinnati, Ohio as: ALWAYS® ALLDAYS® Pantliners with DriWeave® manufactured according to U.S. Patents 4,324,246; 4,463,045; and 6,004,893; ALWAYS® Ultrathin Slender Maxi with Wings manufactured according to U.S. Patents 4,342,314, 4,463,045, 4,556,146, B1 4,589,876, 4,687,478, 4,950,264, 5,009,653, 5,267,992, and Re. 32,649; ALWAYS® Regular Maxi; ALWAYS® Ultra Maxi with Wings; ALWAYS® Maxi with Wings; ALWAYS® Ultra Long Maxi with Wings; ALWAYS® Long Super Maxi with Wings; and ALWAYS® Overnight Maxi with Wings.

Non-limiting examples of interlabial devices which may be provided with a detection means are described in U.S. Patents 5,762,644; 5,885,265; 5,891,126; 5,895,381; 5,916,205; 5,951,537; 5,964,689; 5,968,026; Des. 404,814; and Des. 413,669.

Non-limiting examples of tampons which may be provided with a detection means and applicators therefor, are described in U.S. Patents 4,726,805; 4,846,802; 4,960,417; 5,087,239; 5,279,541; 5,346,468; 5,348,534; 5,531,674; and 5,566,435. In addition, the detection means could also be placed on a digitally insertable tampon.

Blood Detector

The article according to the present invention also includes at least one sensor for the detection of blood. As used herein, the term "sensor" is defined as a component comprising one or more reactive means being adapted to detect one or more target substances (also referred to as analytes) such as microorganisms or related (bio-)molecules (e.g., an enzyme sensor, organella sensor, tissue sensor, microorganism sensor, immunosensor, and chemical or electrochemical sensor), additionally having the capability to provide a signal of

said detection to the wearer, caretaker, or an actuator. When referring to blood herein a typical component of blood, such as hemoglobin or iron, can of course also be used as analyte. The term "reactive" is defined as having the capability to selectively interact with such target substances.

There are 2 categories of sensors which have different sensitivity: biosensors and chemical-/electro-chemical sensors. Generally biosensors function by providing a means of specifically binding, and therefore detecting, a target biologically active analyte. In this way, the biosensor is highly selective, even when presented with a mixture of many chemical and biological entities, such as vaginal discharge. Electrochemical and chemical sensors, on the other hand, which rely on chemically reactive means, generally do not have either the high selectivity or the amplification properties of biosensors but are highly reliable, inexpensive, i.e. useful for commodity products, and often very well established, i.e. proven to be safe for use on human skin. Often the target analyte is a minor component of a complex mixture comprising a multiplicity of biological and other components. Thus, in many biosensor applications, detection of target analytes to the parts-per-billion, parts-per-trillion, or even lower levels is necessary.

According to the present invention an absorbent sanitary article comprises a blood detection means in the form of a specifically selected chemical composition. The chemical composition comprises a color indicating component selected from Gum Guaiac or tetra-methyl-benzidine or combinations thereof which reacts when oxidized and an oxidizing component selected from a peroxide or a per-acid or combinations thereof.

The oxidizing component needs to be stable and insensitive to storage conditions usual for such articles (e.g. in a warm and high humidity environment of a bathroom). If the oxidizing component is a peroxide then physio-chemical stabilization e.g. by use of cyclo-dextrin or metal-chelants is desirable. For per-

acids such stabilization has been found to be less necessary, in particular for the preferred epita-phthalimido-peroxy-hexanoic acid. It is also desirable that the oxidizing component is not activated by the conventional components in sanitary articles and the humidity introduced during wearing (e.g. sweat) or absorption of vaginal discharges without blood component. The same applies of course also to the color indicator.

Not only stabilization of each component but also premature or accidental reaction of the 2 components must be prevented. This can be achieved by separating them physically in that one is applied on one surface of one layer of the sanitary article according to the present invention and the other component is applied on the other side of that layer. Alternatively one component can be applied on the wearer facing surface of one layer and the other component on the wearer facing surface of another layer or the same on the garment facing surface respectively. Finally one component can be applied so that it is located inside a layer e.g. inside the central absorbent core layer while the other component is located on at least part of the surface of that layer. Application of both components on the same layer can be done in an alternate pattern such as stripes, dots or different shapes. The detector components can be provided in discrete parts of a surface of a layer or as full surface coverage of the layer (both components or one alone). The components can be provided either one or both as powder, liquid or as components in an adhesive admixture for better coating.

It is also possible to separate the 2 components by providing at least one, e. g. the oxidizing component, completely separate from the article (and hence from the other component) e. g. in the form of a spray or liquid or powder, which is applied to the article after use to provide the desired detection. Such detection systems could be especially desirable in the context of articles which are worn in close proximity and can be flushed into the toilet (e.g. tampons or interlabial articles). Such systems would address concerns of stability for the separate

component but also of regulatory concerns with materials coming into close contact with the regenerative organs of a woman.

The color indicator and the oxidizing material both are reactive only in aqueous solution. Hence they will only start to react once they have dissolved in the vaginal discharges and are activated by the presence of blood. In order to accelerate and promote the dissolution it is preferred to include a tensioactive material, such as a surfactant, preferably a non-ionic surfactant, or polyethylene-glycol or a combination thereof. The tensioactive should be located such that it is wetted first before the liquid reaches the 2 components.

A highly preferred composition comprises epta-phthalimido-peroxy-hexanoic acid and Gum Guaiac. The Gum Guaiac is a resin found from wood of *Guaiacum officinale* or *Guaiacum santum* (mainly found in Mexico or West India). Gum Guaiac is historically used as flavoring agent for food (and hence has a long record of safe usage by humans) and is known for its ability to indicate the presence of blood or hemoglobin, especially in feces. According to the present invention the Gum Guaiac wood resin is preferred, however one or all of the active components, whether naturally or artificially derived, can be used in stead of the wood resin. The best known active components are guaiacol ($\text{OHC}_6\text{H}_4\text{OCH}_3$, CAS 90-05-1), guaiaconic acid and (furo)-guaiacine.

The components can be applied by coating or spraying provided they are applied in solution. The solvent in this case can either be water or an organic solvent. If desired to remove the solvent then such solvent, especially if it contains water, should evaporate at temperatures below 100 °C.

Additional Detectors

In one particularly preferred embodiment of the present invention the article comprises in addition to the blood detector another separately activated and independently functioning sensor to detect the approach of menstruation.

It is clear that the detection of blood in particular to allow prediction of menstruation cannot be considered to be one hundred percent accurate. First of all, the indicator itself needs to be provided with a high degree of accuracy for only detecting blood, i. e. it should be insensitive to other materials to which it is exposed. On the other hand, the sensitivity should be high to allow detection of low quantities of blood, i.e. prior to its visual recognition. These two competing interests need to be balanced.

Further the presence of (occult) blood is well known as an indication of the approach of menstruation. However also injuries or pathological situations may cause the presence of low levels of blood in vaginal discharges. It is therefore desirable to provide the sanitary article with an additional indicator providing a second signal of the approach of menstruation. Such a signal can be provided, for example, by the change of pH of the vaginal discharge within the days prior to menstruation. In addition or alternatively, hormones such as progesterone or estrogen can be used as additional indicators for the prediction of menstruation.

Most preferred in this context are pH sensors which can also be provided as a color change indicators. pH sensors are well known and are commercially available e.g. from The Merck Company, Darmstadt, Germany. A highly preferred pH sensor, having a well established safety profile in this context is carminic acid, which is used as a food dye, but undergoes a color change in the range between pH 4 and pH 7. Carminic acid is a tricyclic compound which has the compound formula C₂₂-H₂₀-O₁₃.

Regardless of which additional sensors are use if they are provided as color change indicators they preferably can be provided separately, for example in a

different region of the article (of course, such that they will be exposed to vaginal discharges). Alternatively, they can be provided together with the color indicator providing blood detection. In this way the color indicator can be used to provide a multicolor indication.

For example, in the context of Gum Guaiac, which provides a color change to blue and a pH sensor providing e.g. a color change to red, the mixed color sensor (blue and red) would need to turn purple in order to provide an indication with high accuracy and time proximity of the approach of menstruation. A different color change (such as only blue or only red) would indicate that either the menstruation is still some time away or that there is the potential of a health issue (a pH change alone can be indicative - but does not have to - of infections, while blood indication alone can - but does not have to - indicate a wound). For such changes the packaging of the sanitary articles could include an instruction to the user to consider visiting a physician to medically evaluate whether any serious health issue (pathological situation) exists. Of course the absence of a color change would indicate that there is still time until the next menstruation.

In addition to the electrochemical sensors, to supplement the blood detector, the cyclic nature of the hormones of the menstrual cycle (i.e., the full 28 day cycle) make them particularly useful in understanding, the position of an individual during her cycle. This has historically been used in the assessment of fertility, but the use according to the present invention goes beyond current uses of hormones to predict ovulation and pregnancy. For example, progesterone peaks and then drops just prior to menstruation. Estrogen also declines just prior to menstruation. Thus, in combination with other predictors, such as blood detection or pH indication, assay for either of these two hormones will allow reliable prediction of the onset and the presence of menstruation. The timing of the peak of these hormones, along with their subsequent drop, allows a highly accurate marking of the time before menstruation.

The only potential drawback of such hormone sensors is the associated cost and complexity. The lack of simple visualization makes their use in mass produced bulk articles not very attractive. Hence the use of hormone detection in order to predict menstruation is preferably provided in conjunction with blood detection and/or pH detection to improve the accuracy of such predictions in the form of a completely separate detection system.

Use of hormones may also be developed for other points of interest in the cycle. Follicle Stimulating Hormone (FSH) exhibits a peak about one week prior to ovulation, giving more advance timing for pregnancy planning than assays for the luteinizing hormone, which exhibits a sharp peak at the time of ovulation. Two to three days of fertility may be missed when relying only on assay of the luteinizing hormone. Thus, in a diagnostic for ovulation herein, it is highly preferred to measure for both the Follicle Stimulating Hormone and the luteinizing hormone along with estrogen.

A rise in Follicle Stimulating Hormone to a near constant amount signals the approach of menopause. This may be of use in planning healthy approaches to menopause, such as Hormone Replacement Therapy, nutritional changes, and checks for osteoporosis.

The sanitary article according to the present invention can be provided with several additional sensor and detector systems which allow e.g. indications or predictions of health related issues of the individual. Such additional sensor systems are primarily useful for sanitary articles which are used in a care taker situation. In general such additional systems or biosensors comprise a recognition element, or molecular recognition element, that provides detection for a particular analyte. The recognition element may be a biologically derived material such as an enzyme or sequence of enzymes; an antibody; a membrane receptor protein; DNA; an organelle, a natural or synthetic cell membrane; an intact or partial viable or nonviable bacterial, plant or animal cell; or a piece of

plant or mammalian tissues, and generally functions to interact specifically with a target analyte. The recognition element is responsible for the selective recognition of the analyte and creates a physical-chemical signal that provides the basis for the output signal.

Biosensors may include biocatalytic biosensors, and bioaffinity biosensors. In biocatalytic biosensor embodiments, the bio-recognition element is "biocatalytic" and may comprise an enzyme, organelle, piece of plant or mammalian tissue, or whole cells, the selective binding sites "turn over" (i.e., can be used again during the detection process), resulting in a significant amplification of the input signal. Biocatalytic sensors such as these are generally useful for real-time, continuous sensing.

Bioaffinity sensors are generally applicable to bacteria, viruses, and toxins and include chemoreceptor-based biosensors and/or immunological sensors (i.e. immunosensors). Chemoreceptors are complex biomolecular macroassemblies responsible, in part, for a viable organism's ability to sense chemicals in its environment with high selectivity. Chemoreceptor-based biosensors comprise one or more natural or synthetic chemoreceptors associated with a means to provide a signal (visual, electrical, etc.) of the presence or concentration of a target biological analyte. Chemoreceptors may include whole or partial nerve bundles (e.g., from antennae or other sensing organs) and/or whole or partial natural or synthetic cell membranes. On the other hand, the bio-recognition elements of immunosensors are generally antibodies. Antibodies are highly specific and can be made toward bacteria, viruses, fragments of microorganisms (e.g., bacterial cell walls, parasite eggs or portions thereof, etc.), and large biomolecules. Suitable antibodies may be monoclonal or polyclonal. In any case, bioaffinity biosensors are generally irreversible because the receptor sites of the biosensor become saturated when exposed to the target biological analyte. In certain embodiments biocatalytic and bioaffinity biosensors may be combined. Biocatalytic and bioaffinity biosensor systems are described in more detail in

Journal of Chromatography, 510 (1990) 347-354 and in the Kirk-Othmer Encyclopedia of Chemical Technology, 4th ed. (1992), John Wiley & Sons, NY.

The biosensors of the present invention preferably also detect biologically active analytes related to impending (i.e., future presentation of symptoms is likely) or current human systemic disease states, including pathogenic bacteria, parasites, viruses, fungi such as *Candida albicans*, antibodies to pathogens, and/or microbially produced toxins. Additionally, the biosensor may target biologically active analytes related to impending or current localized health issues, such as stress proteins (e.g., cytokines) and IL-1 α (interleukin 1-alpha) that may precede the clinical presentation of skin irritation or inflammation. In preferred embodiments, the biosensor functions as a proactive sensor, detecting and signaling the wearer or caretaker of the impending condition prior to the presentation of clinical symptoms. This allows time to administer prophylactic or remedial treatments to the wearer which can significantly reduce, if not prevent, the severity and duration of the symptoms. Further, the biosensor by detecting the presence of a target biological analyte in the wearer's bodily waste, may detect residual contamination on a surface, such as skin, in contact with the biosensor, and provide an appropriate signal.

The signal generated by the recognition element or elements is communicated visually to the wearer or caretaker, e.g. via a color change visible to the human eye. The signal may be qualitative (e.g., indicating the presence of the target biological analyte) or quantitative (i.e., a measurement of the amount or concentration of the target biological analyte). In any case, the signal is preferably durable i.e., stable and readable over a length of time (typically at least of the same magnitude as the usage period of the article). Further, the sensor may be adapted to detect and/or signal only concentrations of the target biological analyte above or below a predefined threshold level (e.g., in cases wherein the target biological analyte is normally present in the bodily waste).

As described above, the target analyte that the biosensors of the present invention are adapted to detect is pathogenic microorganisms such as the pathogenic microorganisms implicated in human gastrointestinal diseases, especially those resulting in diarrhea. It has been found that severe chronic diarrhea may result in weight loss and permanent physical and mental retardation. A non-limiting list of pathogenic bacteria that the biosensor 60 may detect include any of the various pathogenic strains of *Escherichia coli* (commonly known as *E. Coli*); *Salmonella* strains, including *S. typhi*, *S. paratyphi*, *S. enteritidis*, *S. typhimurium*, and *S. heidelberg*; *Shigella* strains such as *Shigella sonnei*, *Shigella flexneri*, *Shigella boydii*, and *Shigella dysenteriae*; *Vibrio cholerae*; *Mycobacterium tuberculosis*; *Yersinia enterocolitica*; *Aeromonas hydrophila*; *Plesiomonas shigelloides*; *Campylobacter* strains such as *C. jejuni* and *C. coli*; *Bacteroides fragilis*; and *Clostridia* strains, including *C. septicum*, *C. perfringens*, *C. botulinum*, and *C. difficile*. A non-limiting example of a commercially available biosensor adapted to detect *E. coli* is available from AndCare, Inc. of Durham, NC, as test kit #4001. ABTECH, Scientific, Inc., of Yardley, PA offers "bioanalytical biotransducers", available as BB Au-1050.5-FD-X, which may be rendered biospecific (for microorganisms or other target biological analytes as described herein) by covalently immobilizing polypeptides, enzymes, antibodies, or DNA fragments to their surfaces. Other suitable microbial biosensors are described in US Patents 5,869,272 (gram negative organisms); 5,795,717 (*Shigella*); 5,830,341; 5,795,453; 5,354,661; 5,783,399; 5,840,488; 5,827,651; 5,723,330; and 5,496,700.

The target analytes that the biosensors of the present invention are adapted to detect may also be viruses. These may include diarrhea-inducing viruses such as rotavirus, or other viruses such as rhinovirus and human immunodeficiency virus (HIV). An exemplary biosensor adapted to detect HIV is described in US Patents 5,830,341 and 5,795,453, referenced above.

In alternative embodiments, the target analytes that the biosensors of the present invention are adapted to detect may also be parasites, especially those which inhabit the gastrointestinal tract during some point in their life-cycle. Such parasites may include protozoans, worms, and other gastrointestinal parasites. Other examples of parasites which may be detected include *entamoeba histolytica* (which cause amoebic dysentery), *trypana cruzi* (which causes Chagas disease), and *plasmodium falciparum*.

In yet other embodiments, the target analytes the biosensors of the present invention are adapted to detect fungi such as *Candida albicans*. In addition to pathogenic bacteria, certain beneficial colonic bacteria may be detected and/or measured as a health indicator, such as *Bifidobacteria* and *Lactobacillus* strains.

The target analytes that the biosensors of the present invention are adapted to detect may also be proteins or antigens related to skin distress. Preferably, these analytes are detectable on or at the skin surface, preferably prior to the presentation of clinically observable skin irritation. These may include stress proteins such as cytokines, histamine, and other immune response factors including interleukins (such as IL-1 α , IL-2, IL-3, IL-4, and IL-8) and interferons (including interferons α and γ). Again, these are preferably detectable by the biosensor 60 prior to the onset of clinically observable redness, irritation, or dermatitis. Additionally, the biosensors of the present invention may be adapted to detect enzymes, or other biological factors, implicated in skin irritation (e.g., diaper dermatitis), including trypsin, chymotrypsin, and lipase.

The biosensors of the present invention may also comprise bio-recognition systems, including enzymes or binding proteins such as antibodies immobilized onto the surface of physico-chemical transducers. For example, a specific strain of bacteria may be detected via biosensors employing antibodies raised against that bacterial strain. Alternatively, a target bacteria may be detected by a bio-recognition element (including antibodies and synthetic or natural molecular

receptors) specific to extracellular products of the target bacteria, such as toxins produced by that strain (e.g., *E. coli*). Exemplary enzyme electrodes that may be used to detect phenols (e.g. in urine or feces) include tyrosinase based electrodes or polyphenol oxidase enzyme electrodes described in U.S. Patent No. 5,676,820 entitled "Remote Electrochemical Sensor," issued to Joseph Wang et al. on October 14, 1997 and U.S. Patent No. 5,091,299 entitled "An Enzyme Electrode For Use In Organic Solvents," issued to Anthony P. F. Turner et al. on February 25, 1992, respectively

In any of the foregoing examples, the specific microorganism may be directly detected or may be detected by binding a toxin, enzyme, or other protein produced by the organism or an antibody, such as a monoclonal antibody, specific to the organism. Exemplary biosensors adapted to detect proteolytic enzymes described in US Patent 5,607,567 and toxins in US Patents 5,496,452; 5,521,101; and 5,567,301.

The biosensor of the present invention may comprise one or more "proactive sensors". This is especially useful in embodiments where the detection of the target biologically reactive analyte precedes the onset of clinically observable health symptoms. As used in this application, the term "proactive sensor" refers to a sensor that is capable of detecting changes or signals on the body of the wearer (i.e., skin) or in the waste, i.e., inputs, that directly relate or, at a minimum, correlate to the occurrence of an impending or potential health or skin related event. Proactive sensors may respond to one or more specific inputs as described above.

A proactive sensor may detect an impending event or detect a parameter that directly relates, or at a minimum correlates to the occurrence of an impending event, particularly a systemic or skin health event or condition (i.e., the presentation of clinically observable indications or symptoms). An impending event that may be detected or predicted by a proactive sensor of the present

invention may include diarrheal disease, skin irritation or rash (including candidiasis), and/or other types of illness or medical conditions of the wearer such as a parasitic infestation. The detected biological analyte may be one or more steps removed from the actual presentation of clinical symptoms. For example, the biosensor may detect potential precursors to the above conditions (e.g., fecal contamination of the skin that may precede the elicitation of stress proteins which may, in turn, precede clinically observable skin irritation. A parameter that correlates to an event is any measurable input, signal such as one or more of the potential inputs listed above, that correlates with the occurrence of the event within the frame of reference of the system (i.e., a signal caused by the waste or the wearer). Proactive sensors in an article may measure one or more different inputs in order to predict an event. For example, the proactive sensor may monitor for *Candida albicans* in vaginal discharges and residual colonic bacteria on the skin (i.e., detecting residual contamination) both of which are signals that may precede skin irritation.

In biosensor embodiments wherein the bio-recognition element does not produce an easily visible signal (e.g., a color change), the biosensor may include a transducer in communication with the bio-recognition element in order to convert the physico chemical signal from the bio-recognition element into a usable signal to the wearer, caretaker, or component of the article (e.g., and actuator). Exemplary transducers may include electrochemical transducers (including potentiometric, amperometric, and conductimetric transducers), optical transducers (including fluorescence, bioluminescence, total internal reflective resonance, and surface plasmon resonance), thermal transducers, and acoustic transducers, as known in the art. A power source, such as a miniature 3 volt watch battery or printed thin film lithium battery, may be connected with the biosensor to provide any required power.

Examples

As an example a panty liner was provided with a blood indicator and provided indication of the approach of menstruation on different women. The panty liner was a conventional panty liner with a topsheet a secondary topsheet layer between the topsheet and the core, a core layer and a polyethylene film as backsheet.

The blood indicator comprised 2 components, a first component being a composition of 0.2 g of polyethylene-glycol as carrier and tensioactive component with 0.04 g of epta-phtalimido-peroxy-hexanoic acid. The second component was a composition of 0.2 g of polyethylene-glycol as carrier and tensioactive component with 0.02 g of Gum Guaiac. The first component was spread over a 20 mm wide and 130 mm long strip onto the garment facing surface of the secondary topsheet layer. The second component was spread over a 20 mm wide and 130 mm long strip onto the garment facing surface of the absorbent core.

Alternatively a commercially available blood detector strip was inserted into a panty liner as exemplified above between the topsheet and the backsheet also having acceptable results. Such strips are available e.g. as Chroma 6 TM from the Menarini Company, Florence Italy, Arkray Aution Sticks TM, by KDK Corporation of Kyoto, Japan. In another alternative a pH indicator strip was used in the same fashion as the blood indicator strip, again with acceptable results. Such pH indicator strips are available as Spezial Indikator, code 109542, from the Merck Company mentioned above.

CLAIMS

1. Absorbent sanitary article for visual blood detection of a quantity of blood in vaginal discharges deposited on said article which quantity of blood is visually not discernible,

said article is of a layered construction, each layer having a wearer and a garment facing surface and said article comprises

- a liquid permeable topsheet on the wearer facing surface of said article,
- a liquid barrier backsheet on the garment facing surface of said article,
- an absorbent core sandwiched between said liquid permeable topsheet and said liquid barrier backsheet,

said article further comprises a blood detection means, said blood detection means being a chemical composition, said composition comprising a first component selected from a peroxide or a per-acid or combinations thereof, and a second component selected from Gum Guaiac or tetra-methylbenzidine or combinations thereof.

2. Article according to claim 1 characterized in that said composition further comprises
a tensioactive component as dissolving agent, preferably a polyethylene-glycol.
3. Article according to any of the preceding claims characterized in that said first component is epita-phthalimido-peroxy-hexanoic acid and said second component is Gum Guaiac.

4. Article according to any of the preceding claims characterized in that said first and said second components are separated in said absorbent article, preferably one component is provided on the garment facing surface of one layer and the other component is provided on the garment facing surface of a second layer or one component is provided on the wearer facing surface of said core and the other component is provided on the garment facing surface of the other layer.
5. Article according to any of the preceding claims characterized in that said composition is applied to said article in a liquid form, preferably as a solution in an organic carrier or water, said organic carrier being capable of evaporating at temperatures below 100° C after the composition has been applied by spraying or coating.
6. Article according to any of the preceding claims characterized in that said article is a panty liner having a thickness of less than 3 mm.
7. Article according to claim 6 characterized in that said panty liner has a generally triangular shape to be suitable for wearing in a string undergarment.
8. Article according to any of the preceding claims characterized in that said article further comprises additional bio-chemical or electro-chemical or chemical detection means suitable to detect physiological changes in the absorbed vaginal discharges which are indicative of the approach of menstruation, , preferably said additional detection means comprises a pH indicator or a progesterone hormone indicator or an estrogen hormone indicator or combinations thereof.

THIS PAGE BLANK (USPTO)